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## Development of Anti-COVID RNAi Therapeutics Using Human iPSC-Derived Alveolar Epithelial Cells

### Grant Award Details

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Development of Anti-COVID RNAi Therapeutics Using Human iPSC-Derived Alveolar Epithelial Cells

**Grant Type:** Discovery Research Projects

**Grant Number:** DISC1COVID19-12047

**Project Objective:** Synthesis of a Universal Endosomal Escape Domain (uEED) to optimize Anti-COVID siRNA delivery into human iPSC-derived Alveolar Epithelial Cells (AECs).

**Investigator:**

<b>Name:</b>	Steve Dowdy
<b>Institution:</b>	University of California, San Diego
<b>Type:</b>	PI

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**Disease Focus:** COVID-19, Infectious Disease

**Human Stem Cell Use:** iPS Cell

**Award Value:** \$150,000

**Status:** Active

### Grant Application Details

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**Application Title:** Development of Anti-COVID RNAi Therapeutics Using Human iPSC-Derived Alveolar Epithelial Cells

**Public Abstract:****Research Objective**

To optimize a new approach to deliver Anti-COVID siRNAs into human iPSC-derived lung cells that can selectively kill the COVID virus

**Impact**

Our proposal, if successful, will solve the siRNA delivery problem and rapidly open the door to Anti-COVID siRNA therapeutics.

**Major Proposed Activities**

- Complete synthesis of a new delivery device called a Universal Endosomal Escape Domain (uEED)
- Generate a panel of human iPSC-derived lung cells
- Test and optimize the ability of the uEED to deliver Anti-COVID siRNAs into human iPSC-derived lung cells
- Rapidly expand the uEED technology to delivery of Anti-COVID siRNAs in a broader panel of human iPSC-derived lung cells

**Statement of Benefit to California:**

COVID-19 is a deadly health hazard for all Californians, Americans and the world. siRNA-induced RNAi responses are highly selective genetic medicines that have great potential to treat COVID patients and to prophylactically inoculate Californians to prevent their infection. However, due to a delivery problem, we cannot yet deliver siRNAs into lung cells of patients. Our proposal, if successful, will solve the siRNA delivery problem and rapidly open the door to Anti-COVID siRNA therapeutics.

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